

05-22-00

09/554951 PCT

U.S. Application No.
Unknown

International Application No.
PCT/US98/24458

Attorney Docket No.
HOLISED.033APC

Date: May 19, 2000

529 Rec'd PCT/PTO 19 MAY 2000

**TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US)
CONCERNING A FILING UNDER 35 USC 371**

International Application No.: PCT/US98/24458
International Filing Date: November 17, 1998
Priority Date Claimed: November 19, 1997
Title of Invention: USE OF Δ^5 -ANDROSTENE-3 β -OL-7,17-DIONE IN THE TREATMENT OF ARTHRITIS
Applicant(s) for DO/EO/US: Charles E. Weeks

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. (X) This is a **FIRST** submission of items concerning a filing under 35 USC 371.
2. (X) This express request to begin national examination procedures (35 USC 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 USC 371(b) and PCT Articles 22 and 39(1).
3. (X) A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
4. (X) A copy of the International Application as filed (35 USC 371(c)(2))
 - a) () is transmitted herewith (required only if not transmitted by the International Bureau).
 - b) () has been transmitted by the International Bureau.
 - c) (X) is not required, as the application was filed in the United States Receiving Office (RO/US).
5. (X) Amendments to the claims of the International Application under PCT Article 19 (35 USC 371(c)(3))
 - a) () are transmitted herewith (required only if not transmitted by the International Bureau).
 - b) () have been transmitted by the International Bureau.
 - c) () have not been made; however, the time limit for making such amendments has NOT expired.
 - d) (X) have not been made and will not be made.
6. (X) Copies of the International Preliminary Examination Reports dated October 27, 1999 and December 27, 1999 with any annexes thereto, such as any amendments made under PCT Article 34.

Items 11. to 16. below concern other document(s) or information included:

7. (X) International Application as published.
8. (X) PCT Form PCT/IPEA/402.
9. (X) PCT request form.
10. (X) A return prepaid postcard.

U.S. Application No.
Unknown

International Application No.
PCT/US98/24458

Attorney Docket No.
HOLISED.033APC

09/554951

526 Rec'd PCT/PTO

19 MAY 2000

Page 2

Date: May 19, 2000

11. (X) Notification of the Recording of a Change.
12. (X) Request for Change of Applicant Address.
13. (X) The following fees are submitted:

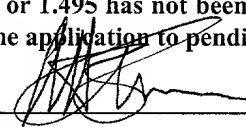
				FEES
BASIC FEE				\$670
CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE	
Total Claims	38 - 20 =	18 ×	\$18	\$324
Independent Claims	2 - 3 =	0 ×	\$78	\$0
Multiple dependent claims(s) (if applicable)			\$260	\$260
TOTAL OF ABOVE CALCULATIONS				\$1,254
Reduction by 1/2 for filing by small entity (if applicable). Verified Small Entity statement must also be filed. (NOTE 37 CFR 1.9, 1.27, 1.28)				\$627
TOTAL NATIONAL FEE				\$627
TOTAL FEES ENCLOSED				\$627
amount to be refunded:				\$0
amount to be charged:				\$0

14. (X) The fee for later submission of the signed oath or declaration set forth in 37 CFR 1.492(e) will be paid upon submission of the declaration.
15. (X) A check in the amount of \$627 to cover the above fees is enclosed.
16. (X) The Commissioner is hereby authorized to charge only those additional fees which may be required, now or in the future, to avoid abandonment of the application, or credit any overpayment to Deposit Account No. 11-1410. A duplicate copy of this sheet is enclosed.

NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.

SEND ALL CORRESPONDENCE TO:

KNOBBE, MARTENS, OLSON & BEAR, LLP
620 Newport Center Drive
Sixteenth Floor
Newport Beach, CA 92660

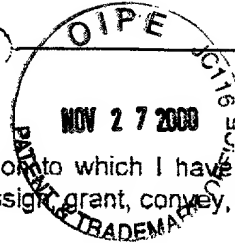

Signature

Ned A. Israelsen
Printed Name

29,655
Registration Number

COPY

VERIFIED STATEMENT (DECLARATION) CLAIMING SMALL ENTITY STATUS (37 CFR 1.9(f) AND 1.27 (c)) - SMALL BUSINESS CONCERN			Docket No. HUM093USPV01
Serial No. 60/066,197	Filing Date November 19, 1997	Patent No.	Issue Date
Applicant/ Weeks Patentee:			
Invention: <u>USE OF ²⁵ ANDROSTENE-3β-OL-7,17-DIONE IN THE TREATMENT OF ARTHRITIS</u>			
<p>I hereby declare that I am:</p> <p><input type="checkbox"/> the owner of the small business concern identified below:</p> <p><input checked="" type="checkbox"/> an official of the small business concern empowered to act on behalf of the concern identified below:</p> <p>NAME OF CONCERN: <u>Humanetics Corporation</u></p> <p>ADDRESS OF CONCERN: <u>600 South Highway 169, Suite 1205, St. Louis Park, Minnesota 55426-1205</u></p> <p>I hereby declare that the above-identified small business concern qualifies as a small business concern as defined in 13 CFR 121.3-18, and reproduced in 37 CFR 1.9(d), for purposes of paying reduced fees under Section 41(a) and (b) of Title 35, United States Code, in that the number of employees of the concern, including those of its affiliates, does not exceed 500 persons. For purposes of this statement, (1) the number of employees of the business concern is the average over the previous fiscal year of the concern of the persons employed on a full-time, part-time or temporary basis during each of the pay periods of the fiscal year, and (2) concerns are affiliates of each other when either, directly or indirectly, one concern controls or has the power to control the other, or a third party or parties controls or has the power to control both.</p> <p>I hereby declare that rights under contract or law have been conveyed to and remain with the small business concern identified above with regard to the above identified invention described in:</p> <p><input type="checkbox"/> the specification filed herewith with title as listed above.</p> <p><input checked="" type="checkbox"/> the application identified above.</p> <p><input type="checkbox"/> the patent identified above.</p> <p>If the rights held by the above-identified small business concern are not exclusive, each individual, concern or organization having rights to the invention is listed on the next page and no rights to the invention are held by any person, other than the inventor, who could not qualify as an independent inventor under 37 CFR 1.9(c) or by any concern which would not qualify as a small business concern under 37 CFR 1.9(d) or a nonprofit organization under 37 CFR 1.9(e).</p>			



Each person, concern or organization to which I have assigned, granted, conveyed, or licensed or am under an obligation under contract or law to assign, grant, convey, or license any rights in the invention is listed below:

- ☒ no such person, concern or organization exists.
☐ each such person, concern or organization is listed below.

FULL NAME

ADDRESS

☐ Individual ☐ Small Business Concern ☐ Nonprofit Organization

FULL NAME

ADDRESS

☐ Individual ☐ Small Business Concern ☐ Nonprofit Organization

FULL NAME

ADDRESS

☐ Individual ☐ Small Business Concern ☐ Nonprofit Organization

FULL NAME

ADDRESS

☐ Individual ☐ Small Business Concern ☐ Nonprofit Organization

Separate verified statements are required from each named person, concern or organization having rights to the invention averring to their status as small entities. (37 CFR 1.27)

I acknowledge the duty to file, in this application or patent, notification of any change in status resulting in loss of entitlement to small entity status prior to paying, or at the time of paying, the earliest of the issue fee or any maintenance fee due after the date on which status as a small entity is no longer appropriate. (37 CFR 1.28(b))

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application, any patent issuing thereon, or any patent to which this verified statement is directed.

NAME OF PERSON SIGNING:

Ronald J. Zenk

TITLE OF PERSON SIGNING

OTHER THAN OWNER:

President

ADDRESS OF PERSON SIGNING:

600 South Highway 169, Suite 1205, St. Louis Park, Minnesota 55426-1205

SIGNATURE:

Ronald J. Zenk

DATE:

1/5/98

USE OF Δ^5 -ANDROSTENE-3 β -OL-7,17-DIONE
IN THE TREATMENT OF
ARTHRITIS

5

FIELD OF THE INVENTION

This invention broadly relates to treatment strategies for arthritis. More specifically, the invention relates to prophylactic, ameliorative and curative drug therapies for arthritis.

10

BACKGROUND

Arthritis is a collective term for number of different conditions that cause pain, swelling and limited movement in joints and connective tissue throughout the body. Specific causes for arthritis are not yet known for most forms of the disease. The condition is usually chronic. The main symptoms of arthritis are joint pain, joint stiffness or inability to move a joint normally, and sometimes swelling that lasts more than two weeks. Most treatment programs include a combination of medication, exercise, rest, use of heat and cold, joint protection techniques, and sometimes surgery.

20

The three most prevalent forms of arthritis are osteoarthritis (OA), fibromyalgia (FM), and rheumatoid arthritis (RA).

25

Osteoarthritis (OA) is a degenerative joint disease characterized by a breakdown of the joint's cartilage. Cartilage functions to cushion the ends of the bones at each joint. A breakdown of the cartilage causes the bones to rub against each other, causing pain and loss of movement. OA primarily affects hands and weight-bearing joints, such as the knee, hips, feet and back. Risk factors for OA include advanced age, obesity, joint injury, and genetic disposition. Suggested causes for OA include an abnormal release of destructive enzymes from the cartilage cells

30

themselves, and inherent defects in the way joints fit together. Most people over 60 suffer from OA, but only about one-third of those over 60 exhibit symptoms of OA.

Conventional treatment of OA focuses on decreasing pain and improving joint movement. Medications include aspirin, acetaminophen, ibuprofen and nonsteroidal anti-inflammatory drugs (herein after NSAIDs) for pain relief and inflammation reduction. Corticosteroid injection directly into affected joints in acute cases is also employed. Other noninvasive techniques commonly used to control OA include heat/cold treatment, exercise, joint protection and weight control.

10

Fibromyalgia (FM) is manifest as widespread pain affecting muscles and attachments to the bone. The patient may also exhibit tender points, specific areas that hurt when pressure is applied. Other symptoms can include fatigue, sleep disturbances, migraine headaches, irritated bowel syndrome, chest pain and nervous system symptoms such as depression.

15

Conventional treatment of FM include use of medications including aspirin, acetaminophen, ibuprofen or NSAIDs for pain relief and inflammation reduction.

20

Rheumatoid arthritis (RA) is an inflammatory disease having many components, including an autoimmune disorder aspect. The autoimmune disorder aspect is generally characterized by inflammation of the membrane lining the joint resulting from an attack upon the joint by the body's own immune system. The inflammation causes pain, stiffness, warmth, redness and swelling. The inflamed joint lining, call the synovium, can invade and damage surrounding bone and cartilage. The involved joint can lose shape and alignment, resulting in pain, loss of movement and possible destruction of the joint. Early in the disease, people may notice general fatigue, soreness, stiffness and aching. Pain usually occurs in the same joints on both sides of the body and will usually start in the hands or feet. RA can also affect wrists, elbows, shoulders, neck, knees, hips and ankles. Other symptoms include lumps, called rheumatoid nodules, under the skin in areas subjected to pressure, such as the back of elbows.

25

30

RA can be diagnosed by a laboratory test for rheumatoid factor, an abnormal substance found in the blood of about 80% of adults with RA. However the presence or absence of rheumatoid factor does not in itself indicate that one has RA. An overall pattern of symptoms, medical history and physical examination are also used in diagnosing RA.

Conventional treatment of RA focuses on reducing swelling, relieving pain and stiffness, and maintaining normal joint function. Medications include NSAIDs for controlling inflammation, joint pain, stiffness and swelling. Disease-modifying drugs include low doses of prednisone, methotrexate, hydroxychloroquine, azulfidine, gold salt and cyclosporin, used alone or in combination. Some combination of exercise, rest, medication, joint protection, physical and occupation therapy, and surgery is also used to treat RA patients.

RA is characterized by striking age-sex disparities. The incidence of RA in women increases steadily from the age of menarche to its maximal incidence around menopause. The disease is uncommon in men under age 45, but its incidence increases rapidly in older men and eventually approaches the incidence in women. These observations strongly suggest that androgens may play some role in RA. Dehydroepiandrosterone (DHEA), an adrenal product, is the major androgen precursor in men and women. Its production is dependent upon age, peaking in the 2nd and 3rd decades in women. DHEA levels are low in both men and women with RA, and recent data show that levels of this hormone may be depressed before onset of the disease. The menopausal peak of RA onset suggests estrogen and/or progesterone deficiency may also play a role in the disease. RA typically remits during pregnancy, in parallel with increasing levels of corticosteroids, estrogens and progesterone. Oral contraceptives, which generate a condition of pseudopregnancy, also decrease the risk of RA. These data argue that adrenal and gonadal steroid hormones affect the development of RA. In addition, several studies indicate that corticosteroid production is inappropriately low in patients with RA.

Traditional treatment regimens for arthritis, including medications for reducing tissue inflammation, often meet with limited success. Hence, the search continues for alternative treatments for arthritis patients.

5

SUMMARY OF THE INVENTION

The invention is directed to the prophylactic, ameliorative and curative treatment of arthritis by administering $\Delta 5$ -androstene- 3β -ol-7,17-dione and precursors thereof which are readily metabolized *in vivo* to $\Delta 5$ -androstene- 3β -ol-7,17-dione but essentially incapable of being metabolized to androgens, estrogens or dehydroepiandrosterone.

15

DETAILED DESCRIPTION OF THE INVENTION INCLUDING A BEST MODE

Arthritis can be treated by administering therapeutic amounts of $\Delta 5$ -androstene- 3β -ol-7,17 dione and precursors thereof which are readily metabolized *in vivo* to $\Delta 5$ -androstene- 3β -ol-7,17-dione but essentially incapable of being metabolized to androgens, estrogens or dehydroepiandrosterone, such as $\Delta 5$ -androstene- 3β -acetoxy-7,17 dione and other 3β esters thereof.

25

Such treatment can be prophylactic, ameliorative or curative in nature.

$\Delta 5$ -Androstene- 3β -ol-7,17-dione

$\Delta 5$ -androstene- 3β -ol-7,17-dione is a derivative of dehydroepiandrosterone (DHEA) which does not appreciably stimulate, increase or otherwise enhance the production of sex hormones. $\Delta 5$ -androstene- 3β -ol-7,17 dione is commercially available from a number of sources including Steraloids, Inc. of Wilton, New Hampshire. A number of procedures are available for synthesizing $\Delta 5$ -androstene- 3β -

ol-7,17 dione from DHEA, with one such procedure described in United States Patent No. 5,296,481.

Precursors of Δ^5 -androstene-3 β -ol-7,17-dione, other than DHEA, may also be usefully employed in the treatment of arthritis. Such precursors are readily metabolized *in vivo* to the active Δ^5 -androstene-3 β -ol-7,17-dione. One example of such a metabolizable precursor is the commercially available Δ^5 -androstene-3 β -acetyl-7,17-dione. The 3 β -acetyl group is readily hydrolyzed *in vivo* by esterases located in the blood and various body tissue to produce the active Δ^5 -androstene-3 β -ol-7,17-dione, and is believed to be less susceptible to oxidation at the 3-position during the manufacturing process relative to Δ^5 -androstene-3 β -ol-7,17-dione.

Administration

Administration Route

The Δ^5 Androstene-3 β -acetoxy-7,17-dione can be administered by virtually any of the commonly accepted practices for the administration of pharmaceutical preparations including specifically, but not exclusively, intravenous injection, mucosal administration, oral consumption, ocular administration, subcutaneous injection, transdermal administration, etc.

Mucosal administration of Δ^5 Androstene-3 β -acetoxy-7,17-dione includes such routes as buccal, endotracheal, inhalation, nasal, pharyngeal, rectal, sublingual, vaginal, etc. For administration through the buccal / inhalation / sublingual / pharyngeal / endotracheal mucosa, the steroid may be formulated as an emulsion, gum, lozenge, spray, tablet or an inclusion complex such as cyclodextrin inclusion complexes. Nasal administration is conveniently conducted through the use of a sniffing powder or nasal spray. For rectal and vaginal administration the steroid may be formulated as a cream, douch, enema or suppository.

Oral consumption of the steroid may be effected by incorporating the steroid into a food or drink, or formulating the steroid into a chewable or swallowable tablet or capsule.

5

Ocular administration may be effected by incorporating the steroid into a solution or suspension adapted for ocular application such as drops or sprays.

Intravenous and subcutaneous administration involves incorporating the steroid into a pharmaceutically acceptable and injectable carrier.

10

For transdermal administration, the steroid may be conveniently incorporated into a lipophilic carrier and formulated as a topical creme or in an adhesive patch.

15

Dose Rate

20

The range of dosages and dose rates effective for achieving the desired biological properties and characteristics may be determined in accordance with standard industry practices. These ranges can be expected to differ depending upon whether the desired response is the prophylactic, ameliorative or curative treatment of arthritis, the specific type of arthritis and the severity of symptoms.

00421756360

EXPERIMENTAL

Experiment 1

(Preparation of Δ^5 Androstene-3 β -acetoxy-7,17-dione)

5

Step One:

(Preparation of Δ^5 Androstene-3-acetoxy-17-one)

A suitable, three-necked, round-bottom flask equipped with an overhead stirrer,
10 reflux condenser, solids addition funnel and 110-volt temperature controller was
charged with a mixture of dichloromethane (90 ml), glacial acetic acid (150 ml), and
acetic anhydride (250 ml). To the mixture was added dehydroepiandrosterone (0.20
moles) purchased from Steraloids, Inc. of Wilton, New Hampshire. The mixture was
stirred to dissolve the solid dehydroepiandrosterone, and anhydrous sodium acetate (35.0
15 g) added. The resulting mixture was heated at 75°C with stirring for 3 hours to
complete the reaction.

The reaction mixture was poured into one liter of water and the resulting slurry
stirred at room temperature for 2 hours. The organic dichloromethane layer was
20 separated from the aqueous layer, and the aqueous layer extracted once with 50 ml of
fresh dichloromethane. The combined organic dichloromethane extract was washed
with water, saturated sodium bicarbonate solution (until neutral), and water. The
resulting washed organic dichloromethane extract was evaporated under reduced
pressure to a volume of 40 ml. Methanol (100 ml) was added to this concentrated extract
25 and the resulting solid mass was cooled at 0°C in a refrigerator for 2 hours.

The resulting solid white product was collected by vacuum filtration on a
Buchner funnel and the filter cake air dried on the funnel to form a first crop of product
weighing 50.5 g. The filtrate mother liquor was concentrated by evaporation under
30 reduced pressure, and cooled at 0°C in a refrigerator. The resulting solid white product
was collected by vacuum filtration on a Buchner funnel and the filter cake air dried on
the funnel to form a second crop of product weighing 10.2 g. The filtrate mother liquor

from the second crop of product was diluted with water and the mixture was cooled at 0°C in a refrigerator. The resulting solid white product was collected by vacuum filtration on a Buchner funnel and the filter cake air dried on the funnel to form a third crop of product weighing 4.2 g.

5

The first, second and third crops of product were combined to produce a total of 64.9 grams of Δ^5 androstene-3-acetoxy-17-one.

10

Theoretical yield	=	66.1 g
First crop yield	=	50.5 g (76.4%)
Second crop yield	=	10.2 g (15.4%)
Third crop yield	=	4.2 g (6.4%)

Step Two:

15

(Preparation of Crude Δ^5 Androstene-3-acetoxy-7,17-dione)

20

A suitable, three-necked, round-bottom flask equipped with an overhead stirrer, reflux condenser, addition funnel, thermometer, mineral oil filled bubbler and a gas inlet tube connected to a nitrogen cylinder, was charged with acetone (3.5 L) and cyclohexane (3.5 L). 1.51 moles of the Δ^5 Androstene-3-acetoxy-17-one prepared in Step One was added to the flask with stirring to dissolved the solid Δ^5 Androstene-3-acetoxy-17-one. 2.48 moles of solid sodium metaperiodate and water (1.1 L) were added to the stirred solution. 14.75 moles of a 70% aqueous solution of *t*-butyl hydroperoxide(2.0 L) was added to the flask through the addition funnel over a one-half hour period.

25

Over the first hour, the reaction mixture temperature rose from 20°C to 32°C. Tap water was added to an external cooling bath and the reaction mixture temperature returned to 20°C. The reaction mixture was constantly vigorously stirred throughout the experiment, and the reaction judged to be complete after 48 hours by TLC monitoring of the disappearance of starting material. The mixture changed from a white slurry to a light yellow slurry over the course of the reaction.

30

The reaction mixture was poured into a stirred ice/water mixture (12 kg ice and 8 L water). Potassium sulfite (3 L of a 45% aqueous solution) was then added to the diluted reaction mixture over 30 minutes (100mL/min) to destroy any remaining oxidant. The diluted mixture was stirred for an additional 2 hours, with ice added as needed to maintain the mixture at 15°C.

The resulting diluted, cooled reaction mixture was transferred to a suitable container and ethyl acetate (3 L) was added to dissolve and extract the product. The resultant mixture was stirred for one-half hour and then allowed to stand so as to permit the organic and aqueous layers to separate. The aqueous layer was examined by TLC, found to contain no product, and discarded. The solids containing organic layer was transferred to a separatory funnel, washed with water (3 x 1.5 L), then washed with a saturated salt solution (1 x 1.5 L). The washed organic layer was dried over sodium sulfate (300 g), with decolorizing carbon (100 g) added. The resulting organic slurry was filtered through a ceramic Buchner funnel containing a 0.5 inch Celite pad (100 g). The filter cake was washed with ethyl acetate (2 x 150 ml) and the washing combined with the filtrate.

The combined organic filtrate was concentrated *in vacuo* to near dryness to produce an off-white semi-solid. The semi-solid was suspended in methanol (400 ml) and again concentrated *in vacuo* to near dryness to produce a semi-solid. The semi-solid was slurried in methanol (600 ml) and the slurry stirred for 2 hours at ambient temperature. The solid product was collected by filtration on a ceramic Buchner funnel, and the solids washed with cold (5°C) methanol (2 x 75 ml). The solid product was dried at 65°C for 48 hours under high vacuum (<1 mm Hg vacuum). The process yielded 232 grams of crude solid Δ^5 Androstene-3-acetoxy-7,17-dione.

Theoretical yield	=	521 g
Actual yield	=	232 g (44.5%)

*Step Three:**(Preparation of Purified Δ^5 Androstene-3-acetoxy-7,17-dione)*METHOD A:

5

A 500 ml round bottom flask equipped with a magnetic stirrer was charged with 25.0 grams of the crude Δ^5 Androstene-3-acetoxy-7,17-dione prepared in Step Two and 300 ml of a mixture of methanol and ethyl acetate (1:1, v/v). The magnetic stirrer was activated and the slurry stirred at room temperature until the crude Δ^5 Androstene-3-acetoxy-7,17-dione was completely dissolved in the solvent mixture to form a first solution. A freshly prepared 10% aqueous solution of sodium bicarbonate (25 ml) was added over 10 minutes to the reaction mixture. The resulting milky mass was stirred at room temperature for 2.5 hours.

15

The reaction mixture was concentrated at room temperature under reduced pressure to 100 ml volume. The concentrated reaction mixture was diluted with 200 ml of ice water and stirred for 30 minutes at 0-5°C. The precipitated solids were collected on a ceramic Buchner funnel, and the aqueous filtrate reserved for additional product recovery. The solids on the funnel were washed with water (until neutral), and methanol (2 x 30 ml), with the methanol washing also reserved for product recovery. The first crop of solids was dried overnight under vacuum to give 18.0 g of purified product.

25

The aqueous filtrate from the first crop of solids was extracted with ethyl acetate (100 ml), and the separated organic extract was washed with water. The solvent of the washed extract was removed under reduced pressure to produce a solid product. This solid product was dissolved in the methanol washing from the first crop of solids, and the solution concentrated to 30 ml volume. Upon cooling the concentrate, a solid precipitate product formed which was collected by vacuum filtration. The second crop of solids was air dried to give 5.2 g of purified product.

30

The mother liquor filtrate from the second crop of solids was diluted with water and cooled. The resulting white solid precipitate was collected by vacuum filtration, and dried overnight at room temperature to give a third crop of 1.0 g of purified product.

- 5 The process yielded a total of 24.2 grams of purified solid Δ^5 Androstene-3-acetoxy-7,17-dione.

Theoretical recovery = 25.0 g

Actual recovery = 24.2 g (96.8%)

10

METHOD B:

- A suitable round bottom flask equipped with a magnetic stirrer was charged with 1.0 gram of the crude Δ^5 Androstene-3-acetoxy-7,17-dione prepared in Step Two and 10 ml of acetone. The magnetic stirrer was activated and the slurry was stirred at room temperature until the crude Δ^5 Androstene-3-acetoxy-7,17-dione was completely dissolved in the acetone. To this solution was added 2.0 g of aluminum oxide basic. The resulting slurry was stirred at room temperature for 1 hour, then filtered through a bed of Celite. The collected solids and Celite bed were washed once with 5 ml of acetone, and the washing combined with the filtrate. The combined filtrate was 20 evaporated to near dryness under reduced pressure to produce a solid product. The solid product was dissolved in a mixture of methanol and isopropyl ether (8:2, v/v) with heating. This solution was cooled at 0-5°C for 30 minutes, resulting in precipitation of a white product. The precipitated solid was collected by vacuum filtration and air dried to 25 give 0.9 grams of purified solid Δ^5 androstene-3-acetoxy-7,17-dione.

Theoretical recovery = 1.0 g

Actual recovery = 0.9 g (90.0%)

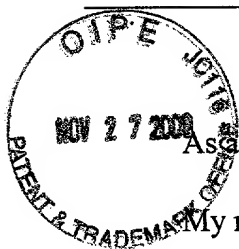
30

I claim:

1. A treatment method comprising treating an arthritis affected patient in need of such treatment with an effective tissue inflammation ameliorative amount of a steroid selected from $\Delta 5$ -androstene- 3β -ol-7,17-dione and metabolizable precursors thereof incapable of being appreciably metabolized to androgens, estrogens or dehydroepiandrosterone.
2. The treatment method of claim 1 wherein the method comprises treating an arthritis affected patient with an effective amount of a tissue inflammation ameliorative amount of a steroid selected from $\Delta 5$ -androstene- 3β -ol-7,17-dione and $\Delta 5$ -androstene- 3β -acetoxy-7,17-dione.
3. The treatment method of claim 1 or 2 wherein the method of treating an arthritis affected patient comprises treating an arthritis affected human patient.
4. The treatment method of claim 3 wherein the method of treating an arthritis affected human patient comprises treating a human patient afflicted with osteoarthritis.
5. The treatment method of claim 3 wherein the method of treating an arthritis affected human patient comprises treating a human patient afflicted with fibromyalgia.
6. The treatment method of claim 3 wherein the method of treating an arthritis affected human patient comprises treating a human patient afflicted with rheumatoid arthritis.
7. The treatment method of claim 3 wherein the method of treating an arthritis affected human patient comprises treating an arthritis affected patient afflicted with arthritis-related tissue inflammation.

8. The treatment method of claim 3 wherein the method of treating an arthritis affected human patient comprises treating a human patient diagnosed with osteoarthritis.
9. The treatment method of claim 3 wherein the method of treating an arthritis affected human patient comprises treating a human patient diagnosed with fibromyalgia.
10. The treatment method of claim 3 wherein the method of treating an arthritis affected human patient comprises treating a human patient diagnosed with rheumatoid arthritis.
11. The treatment method of claim 3 wherein the method of treating an arthritis affected human patient comprises treating an arthritis affected patient diagnosed with arthritis-related tissue inflammation.
12. A treatment method comprising administering an effective arthritis-related tissue inflammation preventative amount of a steroid selected from $\Delta 5$ -androstene- 3β -ol-7,17-dione and metabolizable precursors thereof incapable of being appreciably metabolized to androgens, estrogens or dehydroepiandrosterone to a patient susceptible to arthritis-related tissue inflammation.
13. The treatment method of claim 12 wherein the method comprises treating a patient susceptible to arthritis-related tissue inflammation with an effective arthritis-related tissue inflammation preventative amount of a steroid selected from $\Delta 5$ -androstene- 3β -ol-7,17-dione and $\Delta 5$ -androstene- 3β -acetoxy-7,17-dione.

14. The treatment method of claim 12 or 13 wherein the method of treating a patient susceptible to arthritis-related tissue inflammation comprises treating a human patient susceptible to arthritis-related tissue inflammation.
15. The treatment method of claim 14 wherein the method of treating a patient susceptible to arthritis-related tissue inflammation comprises treating a human patient afflicted with osteoarthritis.
16. The treatment method of claim 14 wherein the method of treating a patient susceptible to arthritis-related tissue inflammation comprises treating a human patient afflicted with fibromyalgia.
17. The treatment method of claim 14 wherein the method of treating a patient susceptible to arthritis-related tissue inflammation comprises treating a human patient afflicted with rheumatoid arthritis.
18. The treatment method of claim 14 wherein the method of treating a patient susceptible to arthritis-related tissue inflammation comprises treating a human patient diagnosed with osteoarthritis.
19. The treatment method of claim 14 wherein the method of treating a patient susceptible to arthritis-related tissue inflammation comprises treating a human patient diagnosed with fibromyalgia.
20. The treatment method of claim 14 wherein the method of treating a patient susceptible to arthritis-related tissue inflammation comprises treating a human patient diagnosed with rheumatoid arthritis.
21. The treatment method of claim 14 wherein the method of treating an arthritis affected human patient comprises treating a human patient diagnosed with osteoarthritis.



DECLARATION - USA PATENT APPLICATION

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name;

I believe I am the original, first and sole inventor of the subject matter which is claimed and for which a patent is sought on the invention entitled USE OF DELTA5-ANDROSTENE-3BETA-OL-7,17-DIONE IN THE TREATMENT OF ARTHRITIS; the specification of which was filed on **May 19, 2000** as Application Serial No. **09/554,951** and was amended on .

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above;

I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, § 1.56;

I hereby claim the benefit under Title 35, United States Codes § 119(e) of any United States provisional application(s) listed below.

Application No.: 60/066,197

Filing Date: November 19, 1997

I hereby claim foreign priority benefits under Title 35, United States Code, § 119(a)-(d) of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed:

PRIOR FOREIGN APPLICATION(S)

Priority
Claimed

No.: **PCT/US98/24458**

Country: **PCT**

Date Filed: **11/17/98**

Yes

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful, false statements may jeopardize the validity of the application or any patent issued thereon.

Inventor's signature

Date _____

Residence: Battle Creek, MI

Citizenship: United States

Post Office Address: 32 Bay Pointe, Battle Creek, Michigan 49015

Send Correspondence To:

KNOBBE, MARTENS, OLSON & BEAR, LLP

Customer No. 20,995

S:\DOCS\NAI\NAI-3246.DOC
080800